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(54) Use of 5-(3,4-dimethoxyphenethyl)methylamino-2-(3,4-dimethoxy-phenyl)-2-isopropylvaleronitrile.

(57) 5-((3,4-dimethoxyphenethyl)methylamino)-2-(3,4-dimethoxy-phenyl)-2-isopropylvaleronitrile or a pharmaceutically acceptable salt thereof is effective to prevent metastasis of cancer.

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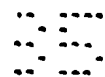
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Use of 5-((3,4-dimethoxyphenethyl)methylamino)-2-
(3,4-dimethoxy-phenyl)-2-isopropylvaleronitrile

This invention relates to a novel agent for preventing metastasis of cancer, i.e. an anti-metastatic agent.

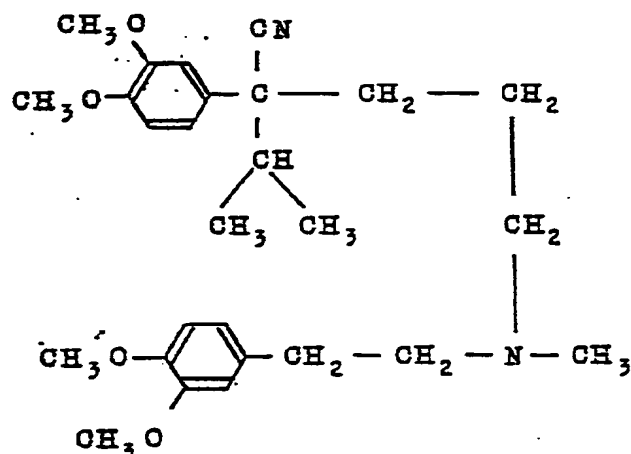
5 More particularly it relates to an anti-metastatic agent comprising 5-[(3,4-dimethoxyphenethyl)methylamino]-2-(3,4-dimethoxyphenyl)-2-isopropyl-



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valeronitrile of the formula

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or a salt thereof as an active ingredient.

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In the recent statistics of the survey, cancer has occupied the first place in death causes in Japan, instead of cerebrovascular diseases. 24% of deaths, i.e. one among four, died of cancer. This mortal disease causes indescribable pain not only to the body but to the mind of a patient. In addition, cancer would most frequently attack those in the prime of life (i.e. in forties to fifties) and playing important roles both in society and in their own homes so that their families also suffer from serious mental and economical damages.

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Therefore various studies to reveal the fundamental cause of cancer and to establish epoch-making processes for the treatment and diagnosis thereof

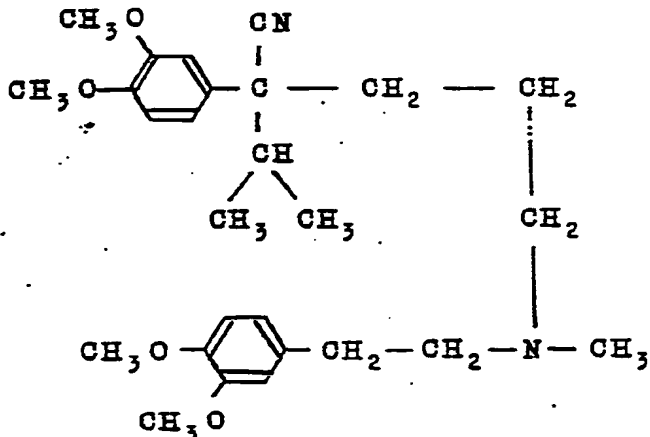
5 have been carried out all over the world to thereby
gain ascendancy over cancer step by step. These
studies have brought about significantly improved
treatments and diagnosis of cancer, so that it can
be completely cured in most cases if detected early
enough.

10 Even if the original tumor is completely removed by
early diagnosis followed by a surgical operation, however,
tumor cells would metastasize to another organ
at the time of the diagnosis in more than half cases.
That is, many patients died of metastasis of cancer.
Accordingly it is one of the most important problems
in the treatment of cancer to prevent its metastasis.

15 Metastasis, which is a specific, complicated
and important characteristic of cancer, would com-
prise many steps such as liberation of cancerous
cells from the primary portion, transfer via blood
or lymph vessels, adhesion to a blood or lymph vessel of
an organ, infiltration and growth. The metastasis of cancer
20 is an important factor governing the recuperation
of a patient. However studies thereon still remain
significantly backward since appropriate experimental
system to evaluate the metastasis is quite limited. The
mechanism of metastasis has not been clarified up and few
25 countermeasures have been established at present.

In order to lower the mortality from cancer, it is a very important problem to prevent and treat the metastasis. Few antimetastatic agents, however, have been known to date.

5 Under these circumstances, we have tried to develop an agent for preventing metastasis of cancer, i.e. an antimetastatic agent, for a long time and found that verapamil, i.e. 5-[(3,4-dimethoxyphenethyl)methyl]-2-(3,4-dimethoxyphenyl)-2-isopropylvaleronitrile of the following formula or its
10 salt such as hydrochloride would be unexpectedly effective as an antimetastatic agent.



Accordingly it is an object of the present invention to provide a novel antimetastatic agent.

25 Verapamil of the above formula has been used as a therapeutic agent for ischemic cardiac diseases

in treating, e.g., stenocardia, coronary arterio-
sclerosis (chronic ischemic cardiac diseases),
silent ischemic cardiac diseases and arteriosclero-
sis cardiac diseases), and myocardial infraction.

5 Verapamil hydrochloride has a melting point
of 138.5 to 140.5°C (decomp.).

To further illustrate the present invention,
the following examples will be given.

10 Example 1

Effect of verapamil on plumonary metastasis of
B16 melanoma BL-6

15 B16 melanoma BL-6, isolated by Dr. Hart
et al. in U.S.A., is a cell line which infiltrates
through a bladder membrane and shows metastatic
potential. 5×10^4 cells of B16 melanoma BL-
6 were inoculated into the tail vein of a male
C57BL/6J mouse. Verapamil hydrochloride was
20 administered intraperitoneally once a day two
days before the inoculation of tumor cells
and three days thereafter, that is,
six times in total. On the 25th day of the trans-
plantation, the mouse was anatomized to observe the
metastasis to the lungs. The degree of the metas-
25 tasis was evaluated by the number of plumonary
nodules. The evaluated values were represented by
range, median and mean + DS. a mark "a"
indicates that a significant difference has
been observed when compared with a control,
30 that is, p is smaller than 0.05 (Student's
t-test). Ten mice were used per a group.
Table 1 shows results.

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Table 1

Dose of verapamil hydrochloride (mg/kg)	No. of pulmonary nodules					
	Median	Range	% to control	Mean \pm SD	% to control	Significant Difference
30	4.5	3~9	32	5.0 \pm 1.7	29	a
40	12	1~19	86	9.8 \pm 6.8	56	a
50	9.5	2~13	68	8.0 \pm 4.3	46	a
Control	14	4~33	100	17.5 \pm 9.3	100	

Example 2: Metastasis of B16 melanoma BL-6 to lungs and lymphonodi

25 x 10⁴ cells of B16 melanoma BL-6 were trans-
planted to the right forefoot of a CS7BL/6J male
5 mouse. Cancerous cells would spantaneously metas-
tasize to the right nodi lymphatic axillares and
lungs with the elapse of time. Verapamil hydro-
chloride was intraperitoneally administered once a
day from the fifth to 16th day (i.e. 11 times) after
10 the transplantation of the cancerous cells. On the
17th day of the transplantation, the right forefoot
including the primary tumor was cut off. On the
38th day the mouse was anatomized to determine the
number of plumonary nodules.

15 Table 2 shows the result.

20

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Table 2

Dose of verapamil hydrochloride (mg/kg)	No. of pulmonary nodules					
	Median	Range	% to control	Mean \pm SD	% to control	Significant Difference
30	11	3~21	147	12.3 \pm 6.5	75	
40	5	1~12 ^a	67	5.8 \pm 3.7	35	
50	1.5	0~12	20	3.5 \pm 4.3	21	a
Control	7.5	1~42	100	16.5 \pm 17.1	100	

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Example 3

Effect of verapamil on plumonary metastasis
of highly metastatic clone NL-17 of mouse colonic
cancer colon 26 adenocarcinome

5 5×10^4 cells of a highly metastatic cell
strain clone NL-17 were transplanted into a vein of
a BALB/C femal mouse. Verapamil hydrochloride was
administered intraperitoneally to the mouse once
a day two days before the inoculation of tumor cells and three
10 days thereafter, i.e. six times in total. On the
23th day of inoculation, the mouse was anatomized
to determine the number of metastatic pulmonary
nodules. Results are shown in Table 3.

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Table 3

Dose of verapamil hydrochloride (mg/kg)	No. of pulmonary nodules					
	Median	Range	% to control	Mean \pm SD	% to control	Significant Difference
60	1	0~96	2.1	13.1 \pm 31.3	15.6	a
75	1.5	0~67	3.2	18.6 \pm 24.5	22.1	a
Control	47.5	5~>200	100	84 \pm 71.3	100	

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Example 4

Effect of verapamil on plumonary metastasis
of highly metastatic clone NL-22 of mouse colonic
cancer colon 26 adnecarcinome

5 1×10^6 cells of a highly metastatic cell
strain NL-22 of mouse colonic cancer colon 26 were
transplanted into the right forefoot of a BALB/C
female mouse. Cancerous cells would spontaneously
metastasize to the lungs with the elapse of time.
10 Verapamil hydrochloride was intraperitoneally
administered to the mouse once a day from the sixth
to 12th day of the transplantation, i.e. six times
in total. On the 13th day of the transplantation,
the right forefoot including the primary carcinoma
15 was cut off. On the 29th day, the mouse was anato-
mized to determine the number of plumonary nodules.

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Table 4

Dose of verapamil hydrochloride (mg/kg)	No. of pulmonary nodules					
	Median	Range	% to control	Mean \pm SD	% to control	Significant Difference
50	34.5	12~55	91	33.1 \pm 14.6	59	
60	22.5	10~41	59	23.5 \pm 8.5	42	a
75	19	7~74	50	27.8 \pm 21.8	50	a
Control	38	22~126	100	55.9 \pm 31.8	100	

Examples 1 to 4 as shown above clearly indicate that the verapamil hydrochloride according to the present invention remarkably prevents metastasis of cancer not only in a single experimental system but
5 also in various experimental systems for cancer metastasis in animals.

Accordingly the verapamil according to the present invention is useful as an excellent agent for preventing metastasis of cancer, i.e. an anti-
10 metastatic agent.

The dose of the verapamil of the present invention as an antimetastatic agent depends on various factors such as the type of cancer and the condition of the patient. It may be usually administered to
15 an adult orally or parenterally in a dose of 10 to 500 mg once to four times a day without any limitation.

It may be formulated into various forms such as powder, grain, granule, tablet, capsule and
20 injection. Formulation may be carried out in a conventional manner with the use of conventional carriers.

In addition to the use as a therapeutic agent administered to cancerous patients, the verapamil
25 of the present invention is further available in

preventing metastasis in those who have received medical treatments such as chemotherapy, endocrinotherapy and immunotherapy, radiotherapy or surgical treatments.

5 Needless to say, the agent of the present invention may be simultaneously administered with other carcinostatic agents.

Toxicity of the verapamil hydrochloride as used in the present invention will now be shown.

10 Acute toxicity

Table 5 shows LD₅₀ (mg/kg) thereof.

Table 5

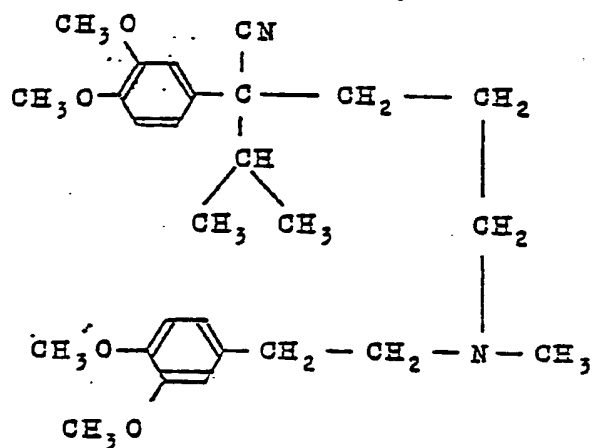
Animal	Sex	Oral	Subcutaneous	Intramuscular	Intravenous
15 Mouse	Male	163	68	-	7.6
Rat	Male	108	107	118	16
	Female	126	-	-	-
Dog	Male and female	>400	-	25	-

20 As described above in detail, the verapamil of the present invention is remarkably effective as an antimetastatic agent. Since metastasis is the cause of deaths due to cancer in most cases, the present invention is extremely valuable.

CLAIM:

Use of 5-((3,4-dimethoxyphenetyl)methylamino)-
2-(3,4-dimethoxy-phenyl)-2-
isopropylvaleronitrile of the formula given
below or a pharmaceutically acceptable salt
5 thereof for the preparing of a composition for pre-
vention of metastasis of cancer.

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